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(21) International Application Number: PCT/US98/00392 (22) International Filing Date: 8 January 1998 (08.01.98) (30) Priority Data: 60/034,194 9 January 1997 (09.01.97) US 60/044,824 25 April 1997 (25.04.97) US 08/924,021 29 August 1997 (29.08.97) US (71) Applicant: ADVANCED CHEMICAL SYSTEMS INTERNATIONAL, INC. [US/US]; 510 Alder Drive, Milpitas, CA 95035 (US). (72) Inventors: WOJTCZAK, William, A.; 350 Elan Village Lane #1, San Jose, CA 95134 (US). GUAN, George; 3900 Moorpark #58, San Jose, CA 95134 (US). FINE, Daniel, N.; 7 Jones Road, Peabody, MA 01960 (US). FINE, Stephen, A.; 7 Jones Road, Peabody, MA 01960 (US). (74) Agents: HAMRICK, Claude, A., S. et al.; Oppenheimer Wolff & Donnelly LLP, Suite 600, Ten Almaden Boulevard, San Jose, CA 95113 (US).		(81) Designated States: CA, ID, IL, JP, KR, SG, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: SEMICONDUCTOR WAFER CLEANING COMPOSITION AND METHOD WITH AQUEOUS AMMONIUM FLUORIDE AND AMINE (57) Abstract <p>A semiconductor wafer cleaning formulation for use in post plasma ashing semiconductor fabrication comprising the following components in the percentage by weight ranges shown: ammonium fluoride and/or a derivative thereof 1-21 %; an organic amine or mixture of two amines 20-55 %; water 23-50 %; a metal chelating agent or mixture of chelating agents 0-21 %.</p>		

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Specification

SEMICONDUCTOR WAFER CLEANING COMPOSITION AND METHOD WITH AQUEOUS AMMONIUM FLUORIDE AND AMINE

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BACKGROUND OF THE INVENTION

Field of the Invention

The present invention relates generally to chemical formulations used in semiconductor wafer fabrication and particularly to chemical formulations that are utilized to remove residue from wafers following a resist plasma ashing step.

Description of the Prior Art

The prior art teaches the utilization of various chemical formulations to remove residue and clean wafers following a resist ashing step. Some of these prior art chemical formulations include alkaline compositions containing amines and/or tetraalkyl ammonium hydroxides, water and/or other solvents, and chelating agents. Still other formulations are based on acidic to neutral solutions containing ammonium fluoride. The various prior art formulations have drawbacks which include unwanted removal of metal or insulator layers and the corrosion of desirable metal layers, particularly aluminum and aluminum-copper alloys and titanium nitride features. There is therefore a need for chemical formulations which effectively remove residue following a resist ashing step which do not attack and potentially degrade delicate structures which are meant to remain on a wafer.

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SUMMARY OF THE INVENTION

A semiconductor wafer cleaning formulation for use in post plasma ashing semiconductor fabrication comprising the following components in the percentage by weight ranges shown:

Ammonium fluoride and/or a derivative thereof;	1-21%
an organic amine or mixture of two amines;	20-55%
water;	23-50%
a metal chelating agent or mixture of chelating agents.	0-21%

It is an advantage of the present invention that it effectively removes inorganic residues following a plasma ashing step.

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It is another advantage of the present invention that it effectively removes metal halide and metal oxide residues following plasma ashing.

It is a further advantage of the present invention that it effectively removes slurry particles of aluminum oxides and other oxides remaining after CMP (chemical mechanical polishing).

It is yet another advantage of the present invention that it provides better stripping performance with less corrosivity than formulations containing ammonium fluoride without amines and amines without ammonium fluoride.

It is yet a further advantage of the present invention that it provides better stripping performance at lower processing temperatures than conventional amine-containing formulations.

It is still a further advantage of the present invention that it includes a chelating agent to prevent metal corrosion and increase stripping effectiveness.

These and other features and advantages of the present invention will become understood to those of ordinary skill in the art upon review of the following detailed description of the preferred embodiments.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention comprises formulations that are suitable for stripping inorganic wafer residues which originate from high density plasma etching followed by ashing with oxygen-containing plasmas. The formulations are also suitable for removing slurry particles of aluminum oxides and other oxides remaining after CMP (chemical mechanical polishing). The formulations contain ammonium fluoride or a derivative of ammonium fluoride, an amine or mixture of amines, water and, optionally, one or more metal chelating agents.

The preferred formulations utilize the following components (percentages are by weight):

Ammonium fluoride and/or a derivative thereof:	1-21%
An organic amine or mixture of two amines	20-55%
Water	23-50%
A Metal chelating agent or mixture of chelating agents:	0-21%

The preferred amines are:

Diglycolamine (DGA)

Methyldiethanolamine (MDEA)

Pentamethyldiethylenetriamine (PMDETA)

Triethanolamine (TEA)

Triethylenediamine (TEDA)

Other amines that are effective include:

5

Hexamethylenetetramine

3, 3-Iminobis (N,N-dimethylpropylamine)

Monoethanolamine

The preferred fluoride sources are:

10

Ammonium fluoride

Triethanolammonium fluoride (TEAF)

Other fluoride sources that are effective include:

Diglycolammonium fluoride (DGAF)

15

Tetramethylammonium fluoride (TMAF)

Triethylamine tris (hydrogen fluoride) (TREAT-HF)

The metal chelating agents that are effective include:

Acetoacetamide

20

Ammonium carbamate

Ammonium pyrrolidinedithiocarbamate (APDC)

Dimethyl malonate

Methyl acetoacetate

N-Methyl acetoacetamide

25

2,4-Pentanedione

Tetramethylammonium thiobenzoate

Tetramethylammonium trifluoroacetate

Tetramethylthiuram disulfide (TMTDS)

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The combination of ammonium fluoride or a substituted fluoride source as identified above with an amine (other than as a surfactant in an amount of 1% or less) provides better stripping performance with less corrosivity than formulations containing ammonium fluoride without amines and formulations containing amines without ammonium fluoride. In addition, the resulting alkaline solutions were effective at lower processing temperatures (21°-40° C)

35 than conventional amine-containing formulations.

The use of 1,3-dicarbonyl compounds as chelating agents and to prevent metal corrosion also appears to increase effectiveness.

In the prior art amines are limited to 1% or less of the formulation and serve only as surfactants, or amines are not utilized as formulation ingredients at all. Also, the prior art formulations are acidic (pH < 7). In the formulations of the present invention the amines are present as major components and play major roles in stripping, and the formulations are basic (pH > 7).

Several representative examples of preferred formulations are:

	Triethanolamine	45%
10	Ammonium fluoride	5%
	Water	50%
	Diglycolamine	55%
	Ammonium fluoride	5%
15	2,4-Pentanedione	10%
	Water	30%
	Triethanolamine	27.1%
	TEAF	20.3%
20	2,4-Pentanedione	10%
	Water	42.6%
	PMDETA	45%
	Ammonium fluoride	5%
25	Methyl acetoacetate	6%
	Water	44%
	PMDETA	45%
	Ammonium fluoride	1%
30	2,4-Pentanedione	8%
	APDC	15%
	Water	31%
	PMDETA	55%
35	Ammonium fluoride	1%
	Dimethyl malonate	13.2%

	TMTDS	6%
	Water	24.8%
	TEA	36%
5	PMDETA	16%
	Ammonium fluoride	12%
	Acetoacetamide	10%
	Water	28%
10	TEA	45%
	Ammonium fluoride	11.4%
	Tetramethylammonium-trifluoroacetate	17%
	Water	27%
15	Triethanolamine	45-52%
	Ammonium fluoride	3-10%
	2,4-Pentanedione	5-10%
	Water	35-44%
20	PMDETA	38-45%
	Ammonium fluoride	5%
	2,4-Pentanedione	10%
	Water	40-47%
25	PMDETA	38%
	TMAF	5%
	Ammonium fluoride	2%
	2,4-Pentanedione	10%
	Water	45%
30	PMDETA	38%
	Ammonium fluoride	1%
	2,4-Pentanedione	10.7%
	Ammonium carbamate	10%
35	Water	38.3%

The inventors expect that closely related ingredients would be expected to show comparable performance to those utilized in our formulations.

These include:

A. Other organic amines are expected to be suitable.

5 B. Other substituted ammonium fluorides are expected to be suitable. These would have the general formula, $R_1R_2R_3R_4NF$ in which the R groups are Hydrogen atoms and/or aliphatic groups.

C. Other metal chelating agents including:

10 1. Other 1,3-dicarbonyl compounds are expected to display comparable performance. These would have the following general structure:

$X-CHR-Y$ in which

R is either a Hydrogen atom or an aliphatic group and

15 X and Y are functional groups containing multiply bonded moieties known to have electron-withdrawing properties, for example X and Y may be $CONH_2$, $CONHR'$, CN , NO_2 , SOR' , SO_2Z in which R' represents an alkyl group and Z represents another atom or group. X and Y may be identical or different.

2. Other amine trifluoroacetates are also expected to be suitable as chelating agents. These would have the general formula, $R_1R_2R_3R_4N^+ O_2CCF_3$ in which the R groups are Hydrogen atoms and/or aliphatic groups.

20 D. It would also be expected that inclusion of optional components such as surfactants, stabilizers, corrosion inhibitors, buffering agents, and cosolvents would constitute obvious additions to those practiced in the art.

The formulations of the present invention are particularly useful on wafers which have been etched with chlorine- or fluorine-containing plasmas followed by oxygen plasma ashing.
25 The residues generated by this type of processing typically contain inorganic materials such as, but not limited to, aluminum oxide, titanium oxide and aluminum fluoride. These residues are often difficult to dissolve completely without causing corrosion of metal and titanium nitride features required for effective device performance. Also, metal oxide and silicon oxide slurry particles remaining after CMP will also be effectively removed by these formulations.

30 Three types of commercially generated wafers containing via structures were evaluated using the formulations of the present invention. In each case, following plasma etching and ashing, immersion of wafers in the formulation of the present invention was performed in baths at 21-60°C for 10-30 minutes (preferred: 21-35°C for 20-30 minutes) followed by washing with deionized water and drying with a stream of nitrogen gas. It is expected that the formulations
35 can also be applied by spraying onto the wafers in an automated spray tool followed by a water rinse.

The three via structures were:

1. 0.8 micron diameter, four layer vias comprised of silicon oxide top and second layers, a third layer of titanium nitride, and a bottom layer of aluminum, silicon, copper (AlSiCu) alloy. The substrate was silicon oxide.

5 2. One micron diameter, two-layer vias comprised of a top layer of silicon oxide (7000 Angs. thick) and a bottom layer of titanium nitride (1200 Angs. thick) on top of a silicon substrate.

10 3. 1.6 micron wide, 4-layer vias with a top layer of titanium nitride (40 Nm. thick), a second layer of silicon oxide (1.3 microns thick), and a bottom layer of aluminum/copper on a silicon oxide substrate.

The present formulations were rated according to relative stripping effectiveness and corrosivity of the formulations. The preferred formulations scored best and, in overall performance based on both stripping effectiveness and low corrosivity, are approximately equal.

15 While the present invention has been described with reference to certain preferred embodiments, it will be understood by those skilled in the art that various alterations and modifications may be made therein without departing from the true spirit and scope of the invention. It is therefore intended that the following claims cover all such alterations and modifications which nevertheless include the true spirit and scope of the invention.

20 What I claim is:

CLAIMS

1. A semiconductor wafer cleaning formulation for use in post plasma ashing semiconductor fabrication comprising the following components in the percentage by weight ranges shown:

	a fluoride source;	1-21%
5	at least one organic amine;	20-55%
	water;	23-50%
	at least one metal chelating agent.	0-21%.

2. A cleaning formulation as described in claim 1 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride; and
Triethanolammonium fluoride (TEAF).

3. A cleaning formulation as described in claim 1 wherein said organic amine is chose from the group consisting of:

Diglycolamine (DGA),
Methyldiethanolamine (MDEA),
5 Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA), and
Triethylenediamine (TEDA).

4. A cleaning formulation as described in claim 1 wherein said metal chelating agent is chosen from the group consisting of:

Acetoacetamide,
Ammonium carbamate,
5 Ammonium pyrrolidinedithiocarbamate (APDC),
Dimethyl malonate,
Methyl acetoacetate,
N-Methyl acetoacetamide,
2,4-Pentanedione,
10 Tetramethylammonium thiobenzoate,
Tetramethylammonium trifluoroacetate, and
Tetramethylthiuram disulfide (TMTDS).

5. A cleaning formulation as described in claim 1 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride,
Triethanolammonium fluoride (TEAF),
5 Diglycolammonium fluoride (DGAF),
Tetramethylammonium fluoride (TMAF), and
Triethylamine tris (hydrogen fluoride) (TREAT-HF).

6. A cleaning formulation as described in claim 1 wherein said organic amine is chosen from the group consisting of:

Diglycolamine (DGA),
Methyldiethanolamine (MDEA),
5 Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA),
Triethylenediamine (TEDA),
Hexamethylenetetramine,
3, 3-Iminobis (N,N-dimethylpropylamine), and
10 Monoethanolamine.

7. A cleaning formulation as described in claim 1 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride,
Triethanolammonium fluoride (TEAF),
5 Diglycolammonium fluoride (DGAF),
Tetramethylammonium fluoride (TMAF), and
Triethylamine tris (hydrogen fluoride) (TREAT-HF);
said organic amine is chosen from the group consisting of:

Diglycolamine (DGA),
10 Methyldiethanolamine (MDEA),
Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA),
Triethylenediamine (TEDA),
Hexamethylenetetramine,
15 3, 3-Iminobis (N,N-dimethylpropylamine), and
Monoethanolamine;

and said metal chelating agent is chosen from the group consisting of:

Acetoacetamide,
 Ammonium carbamate,
 Ammonium pyrrolidinedithiocarbamate (APDC),
 Dimethyl malonate,
 5 Methyl acetoacetate,
 N-Methyl acetoacetamide,
 2,4-Pentanedione,
 Tetramethylammonium thiobenzoate,
 Tetramethylammonium trifluoroacetate, and
 10 Tetramethylthiuram disulfide (TMTDS).

8. A cleaning formulation as described in claim 1 wherein said fluoride source has the general formula $R_1R_2R_3R_4NF$ in which the R groups are hydrogen atoms and/or aliphatic groups, and wherein said metal chelating agent would have the following general structure:

X-CHR-Y in which

5 R is either a hydrogen atom or an aliphatic group and

X and Y are functional groups containing multiply bonded moieties known to have electron-withdrawing properties, wherein X and Y may be $CONH_2$, $CONHR'$, CN , NO_2 , SOR' , SO_2Z in which R' represents an alkyl group and Z represents another atom or group, and wherein X and Y may be identical or different.

9. A cleaning formulation as described in claim 1 wherein said fluoride source⁴ has the general formula $R_1R_2R_3R_4NF$ in which the R groups are hydrogen atoms and/or aliphatic groups, and wherein said metal chelating agent would have the general formula, $R_1R_2R_3R_4N^{+}O_2CCF_3$ in which the R groups are hydrogen atoms and/or aliphatic groups.

10. A cleaning formulation as described in claim 1 being comprised of:

Triethanolamine	45%
Ammonium fluoride	5%
Water	50%.

11. A cleaning formulation as described in claim 1 being comprised of:

Diglycolamine	55%
Ammonium fluoride	5%
2,4-Pentanedione	10%
5 Water	30%.

12. A cleaning formulation as described in claim 1 being comprised of:

Triethanolamine	27.1%
TEAF	20.3%
2,4-Pentanedione	10%
5 Water	42.6%.

13. A cleaning formulation as described in claim 1 being comprised of:

PMDETA	45%
Ammonium fluoride	5%
Methyl acetoacetate	6%
5 Water	44%.

14. A cleaning formulation as described in claim 1 being comprised of:

PMDETA	45%
Ammonium fluoride	1%
2,4-Pentanedione	8%
5 APDC	15%
Water	31%.

15. A cleaning formulation as described in claim 1 being comprised of:

PMDETA	55%
Ammonium fluoride	1%
Dimethyl malonate	13.2%
5 TMTDS	6%
Water	24.8%.

16. A cleaning formulation as described in claim 1 being comprised of:

TEA	36%
PMDETA	16%
Ammonium fluoride	12%
5 Acetoacetamide	10%
Water	28%.

17. A cleaning formulation as described in claim 1 being comprised of:

TEA	45%
Ammonium fluoride	11.4%

Tetramethylammonium-trifluoroacetate	17%
Water	27%.

18. A cleaning formulation as described in claim 1 being comprised of:

Triethanolamine	45-52%
Ammonium fluoride	3-10%
2,4-Pentanedione	5-10%
Water	35-44%.

19. A cleaning formulation as described in claim 1 being comprised of:

PMDETA	38-45%
Ammonium fluoride	5%
2,4-Pentanedione	10%
Water	40-47%.

20. A cleaning formulation as described in claim 1 being comprised of:

PMDETA	38%
TMAF	5%
Ammonium fluoride	2%
2,4-Pentanedione	10%
Water	45%.

21. A cleaning formulation as described in claim 1 being comprised of:

PMDETA	38%
Ammonium fluoride	1%
2,4-Pentanedione	10.7%
Ammonium carbamate	10%
Water	38.3%.

22. A method for fabricating a semiconductor wafer including the steps comprising:

plasma etching a metalized layer from a surface of the wafer;
plasma ashing a resist from the surface of the wafer following the metal etching step;
cleaning the wafer in a following step using a chemical formulation including the

following components in the percentage by weight ranges shown:

a fluoride source;	1-21%
at least one organic amine;	20-55%

water;

23-50%

at least one metal chelating agent.

0-21%.

23. A cleaning formulation as described in claim 22 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride; and

Triethanolammonium fluoride (TEAF).

24. A method as described in claim 22 wherein said organic amine is chose from the group consisting of:

Diglycolamine (DGA),

Methyldiethanolamine (MDEA),

5 Pentamethyldiethylenetriamine (PMDETA),

Triethanolamine (TEA), and

Triethylenediamine (TEDA).

25. A method as described in claim 22 wherein said metal chelating agent is chosen from the group consisting of:

Acetoacetamide,

Ammonium carbamate,

5 Ammonium pyrrolidinedithiocarbamate (APDC),

Dimethyl malonate,

Methyl acetoacetate,

N-Methyl acetoacetamide,

2,4-Pentanedione,

10 Tetramethylammonium thiobenzoate,

Tetramethylammonium trifluoroacetate, and

Tetramethylthiuram disulfide (TMTDS).

26. A method as described in claim 22 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride,

Triethanolammonium fluoride (TEAF),

5 Diglycolammonium fluoride (DGAF),

Tetramethylammonium fluoride (TMAF), and

Triethylamine tris (hydrogen fluoride) (TREAT-HF).

27. A method as described in claim 22 wherein said organic amine is chosen from the group consisting of:

Diglycolamine (DGA),
Methyldiethanolamine (MDEA),
5 Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA),
Triethylenediamine (TEDA),
Hexamethylenetetramine,
3, 3-Iminobis (N,N-dimethylpropylamine), and
10 Monoethanolamine.

28. A method as described in claim 22 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride,
Triethanolammonium fluoride (TEAF),
5 Diglycolammonium fluoride (DGAF),
Tetramethylammonium fluoride (TMAF), and
Triethylamine tris (hydrogen fluoride) (TREAT-HF);
said organic amine is chosen from the group consisting of:

Diglycolamine (DGA),
10 Methyldiethanolamine (MDEA),
Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA),
Triethylenediamine (TEDA),
Hexamethylenetetramine,
15 3, 3-Iminobis (N,N-dimethylpropylamine), and
Monoethanolamine;

and said metal chelating agent is chosen from the group consisting of:

Acetoacetamide,
Ammonium carbamate,
20 Ammonium pyrrolidinedithiocarbamate (APDC),
Dimethyl malonate,
Methyl acetoacetate,
N-Methyl acetoacetamide,
2,4-Pentanedione,
25 Tetramethylammonium thiobenzoate,

Tetramethylammonium trifluoroacetate, and
Tetramethylthiuram disulfide (TMTDS).

29. A method as described in claim 22 wherein said fluoride source has the general formula $R_1R_2R_3R_4NF$ in which the R groups are hydrogen atoms and/or aliphatic groups, and wherein said metal chelating agent would have the following general structure:

X-CHR-Y in which

5 R is either a hydrogen atom or an aliphatic group and

X and Y are functional groups containing multiply bonded moieties known to have electron-withdrawing properties, wherein X and Y may be $CONH_2$, $CONHR'$, CN , NO_2 , SOR' , SO_2Z in which R' represents an alkyl group and Z represents another atom or group, and wherein X and Y may be identical or different.

30. A method as described in claim 22 wherein said fluoride source has the general formula $R_1R_2R_3R_4NF$ in which the R groups are hydrogen atoms and/or aliphatic groups, and wherein said metal chelating agent would have the general formula, $R_1R_2R_3R_4N^+ \cdot O_2CCF_3$ in which the R groups are hydrogen atoms and/or aliphatic groups.

31. A method for fabricating a semiconductor wafer including the steps comprising:
plasma etching a metalized layer from a surface of the wafer;
plasma ashing a resist from the surface of the wafer following the metal etching step;
cleaning the wafer in a following step using a chemical formulation including the

5 following components in the percentage by weight ranges shown:

a fluoride source;	1-21%
at least one organic amine;	20-55%
water;	23-50%
at least one metal chelating agent.	0-21%.

32. A cleaning formulation as described in claim 31 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride; and
Triethanolammonium fluoride (TEAF).

33. A method as described in claim 31 wherein said organic amine is chose from the group consisting of:

Diglycolamine (DGA),

Methyldiethanolamine (MDEA),
5 Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA), and
Triethylenediamine (TEDA).

34. A method as described in claim 31 wherein said metal chelating agent is chosen from the group consisting of:

Acetoacetamide,
Ammonium carbamate,
5 Ammonium pyrrolidinedithiocarbamate (APDC),
Dimethyl malonate,
Methyl acetoacetate,
N-Methyl acetoacetamide,
2,4-Pentanedione,
10 Tetramethylammonium thiobenzoate,
Tetramethylammonium trifluoroacetate, and
Tetramethylthiuram disulfide (TMTDS).

35. A method as described in claim 31 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride,
Triethanolammonium fluoride (TEAF),
5 Diglycolammonium fluoride (DGAF),
Tetramethylammonium fluoride (TMAF), and
Triethylamine tris (hydrogen fluoride) (TREAT-HF).

36. A method as described in claim 31 wherein said organic amine is chosen from the group consisting of:

Diglycolamine (DGA),
Methyldiethanolamine (MDEA),
5 Pentamethyldiethylenetriamine (PMDETA),
Triethanolamine (TEA),
Triethylenediamine (TEDA),
Hexamethylenetetramine,
3, 3-Iminobis (N,N-dimethylpropylamine), and
10 Monoethanolamine.

37. A method as described in claim 31 wherein said fluoride source is chosen from the group consisting of:

Ammonium fluoride,

Triethanolammonium fluoride (TEAF),

5 Diglycolammonium fluoride (DGAF),

Tetramethylammonium fluoride (TMAF), and

Triethylamine tris (hydrogen fluoride) (TREAT-HF);

said organic amine is chosen from the group consisting of:

Diglycolamine (DGA),

10 Methyl-diethanolamine (MDEA),

Pentamethyl-diethylenetriamine (PMDETA),

Triethanolamine (TEA),

Triethylenediamine (TEDA),

Hexamethylenetetramine,

15 3, 3-Iminobis (N,N-dimethylpropylamine), and

Monoethanolamine;

and said metal chelating agent is chosen from the group consisting of:

Acetoacetamide,

Ammonium carbamate,

20 Ammonium pyrrolidinedithiocarbamate (APDC),

Dimethyl malonate,

Methyl acetoacetate,

N-Methyl acetoacetamide,

2,4-Pentanedione,

25 Tetramethylammonium thiobenzoate,

Tetramethylammonium trifluoroacetate, and

Tetramethylthiuram disulfide (TMTDS).

38. A method as described in claim 31 wherein said fluoride source has the general formula $R_1R_2R_3R_4NF$ in which the R groups are hydrogen atoms and/or aliphatic groups, and wherein said metal chelating agent would have the following general structure:

X-CHR-Y in which

5 R is either a hydrogen atom or an aliphatic group and

X and Y are functional groups containing multiply bonded moieties known to have electron-withdrawing properties, wherein X and Y may be $CONH_2$, $CONHR'$, CN , NO_2 , SOR' ,

SO₂Z in which R' represents an alkyl group and Z represents another atom or group, and wherein X and Y may be identical or different.

39. A method as described in claim 31 wherein said fluoride source has the general formula R₁R₂R₃R₄NF in which the R groups are hydrogen atoms and/or aliphatic groups, and wherein said metal chelating agent would have the general formula, R₁R₂R₃R₄N⁺ O₂CCF₃ in which the R groups are hydrogen atoms and/or aliphatic groups.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US98/00392

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : C11D 3/24, 3/30; B08B 3/08

US CL : 510/175, 176; 134/1.3, 29, 38, 42

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 510/175, 176; 134/1.3, 29, 38, 42

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
None

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, CAS ONLINE

search terms: ammonium fluoride, triethanolamine, hydrofluoric, hydrogen fluoride, triethanolammonium fluoride

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E, Y	US 5,709,756 A (WARD et al.) 20 January 1998; column 2, line 1, through column 3, line 23.	1, 8, 9, 22, 29, 30, 31, 38, 39
Y	NOGUCHI, H, et al., Kinki Aruminyumu Hyomen Shori Kenkyukai Kaishi, 1996, 179, 3-8. See Abstract.	1-10, 22-32, 37-39
P, A	US 5,698,503 A (WARD et al.) 16 December 1997.	
A	US 5,571,447 A (WARD et al.) 05 November 1996.	
A	US 5,560,857 A (SAKON et al.) 01 October 1996.	

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
B earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A* document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means	
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 11 MARCH 1998	Date of mailing of the international search report 17 APR 1998
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer DOUGLAS J. MCGINTY Telephone No. (703) 308-0661

INTERNATIONAL SEARCH REPORTInternational application No.
PCT/US98/00392**C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4,863,563 A (SCARDERA et al.) 05 September 1989.	
A	US 4,165,295 A (VANDER MEY) 21 August 1979.	
A	US 4,215,005 A (VANDER MEY) 29 July 1980.	
A	US 4,113,551 A (BASSOUS et al.) 12 September 1978.	
P, A	EP 812,011 A (WAKO PURE CHEMICAL INDUSTRIES) 10 December 1997.	
A	EP 680,078 A (MITSUBISHI GAS CHEMICAL COMPANY, INC.) 02 November 1995.	
A	GB 2,287,827 A (NEC CORPORATION) 27 September 1995.	
A	JP 8,250,461 A (KANTO KAGAKU) 27 September 1996.	
A	DD 153,308 A (GERMAN DEMOCRATIC REPUBLIC) 30 December 1981.	